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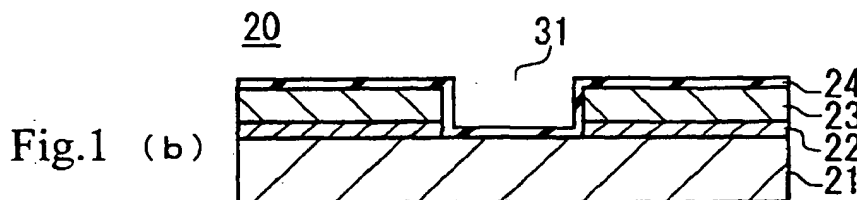
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(54) **Process for producing barrier film**

(57) A thin nitride film having a low resistance is formed at a low film-forming temperature.

In the step of forming a thin nitride film 24 of a high temperature-melting point metal by introducing a feed-stock gas having the high temperature-melting point metal and a reductive nitrogen-containing gas having a nitrogen atom into a vacuum atmosphere, an auxiliary

reductive gas free from nitrogen is also introduced. The high temperature-melting point metal deposited due to the auxiliary reductive gas compensates for the deficiency of the high temperature-melting point metal of the deposited nitride and thus enable the growth of the thin nitride film 24 having a low resistance.



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Description

FIELD OF THE INVENTION

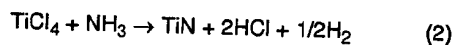
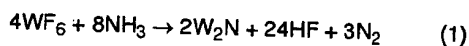
[0001] This invention relates to the technical field of metal interconnecting for semiconductor devices. More particularly, it relates to a process for producing a barrier film which is to be located between a copper interconnecting film and an insulation film, a barrier film which is to be located between a film containing Si, GaAs, etc. and a metal interconnecting for preventing silicidation, and a barrier film which is to be located between a highly dielectric film or a ferroelectric film and an electrode.

BACKGROUND OF THE INVENTION

[0002] In recent years, it has been required to more and more speed up the operations of semiconductor devices. To satisfy this requirement, studies have been undertaken on low-resistant copper interconnectings as a substitute for aluminum interconnectings.

[0003] However, copper occurs as an impurity in semiconductor crystals. In addition, it suffers from a problem of having a large diffusion coefficient in silicon crystals or silicon oxide. Therefore, it has been a practice that a thin film comprising the nitride of a high temperature-melting point metal (a thin tungsten nitride film, etc.) is formed as a barrier film on the surface of a silicon substrate or a thin silicon oxide film and then a copper interconnecting film is formed on the surface of the barrier film.

[0004] To form such a barrier film, the sputtering method, the heat CVD method or the PE-CVD method is used. In the sputtering method, a high temperature-melting point metal is employed as a target. In the heat CVD method, a thin nitride film is formed by the following reduction reactions. Formula (1) shows a case wherein tungsten is used, while formula (2) shows another case wherein titanium is used.



[0005] In case of forming a semiconductor device with multi-layered interconnecting, it is needed to laminate copper interconnectings while inserting interlayer insulation films between them. In a semiconductor device to be operated at a high speed, the resistance of the copper interconnectings as well as the capacity of the interlayer insulation films and the resistance of the barrier films should be lowered so as to minimize signal transfer delay. More concretely speaking, a barrier film should have a resistance as low as 200 to 300 $\mu\Omega\text{cm}$.

[0006] Although a thin nitride film having a low resistance can be formed by the sputtering method, only poor step coverage can be achieved thereby. Thus,

no uniform barrier film can be formed in a viahole with a high aspect ratio by this method.

[0007] By the heat CVD method, on the other hand, a uniform barrier thin film can be formed in a viahole. However, the upper limit of the film-forming temperature in the heat CVD method resides in 400 to 500°C, since the dielectric constants of interlayer insulation films with low dielectric constants would be increased when exposed to a high temperature exceeding 500°C. At such a low film-forming temperature, the resistivity of, for example, a thin tungsten nitride film attains several thousand $\mu\Omega\text{cm}$, which makes it impossible to give barrier films with low resistance.

[0008] With respect to the CVD methods, barrier films with low resistance can be formed at a low temperature by the MOCVD method with the use of organometallic compounds or the plasma CVD method. However, the organometallic compounds are expensive, while the plasma CVD method has a problem of achieving only poor step coverage. Thus, these methods are not usable in practice.

SUMMARY OF THE INVENTION

[0009] The present invention, which has been made to overcome the above-described troubles encountering in the conventional art, aims at providing a barrier film having a value of low resistivity and good step coverage.

[0010] The present inventors analyzed thin films of high temperature-melting point metal nitrides formed by the conventional heat CVD method and, as a result, found that the high temperature-melting point metal atoms were provided only in an insufficient amount. In the case of tungsten, for example, a tungsten nitride in the conventional art fails to establish the stoichiometric composition (W_2N) but shows a composition W_xN wherein x ranges from about 1.5 to 1.6. It is assumed that such insufficient supply of metal atoms in a nitride might worsen the crystallinity of the thin nitride film, thereby elevating the value of the resistance.

[0011] In the present invention which has been completed based on the above-described finding, attempts are made to approximate the composition of the nitride of a high temperature-melting point metal closely to the stoichiometric level. To achieve this object, the present invention relates to a process for producing a barrier film which comprises the steps of providing a substrate in a vacuum atmosphere, introducing a feed-stock gas having a high temperature-melting point metal in its structure and a reductive nitrogen-containing gas having a nitrogen atom into said vacuum atmosphere, and forming a thin film of the nitride of said high temperature-melting point metal on said substrate, wherein a nitrogen-free auxiliary reductive gas is introduced into said vacuum atmosphere.

[0012] The present invention relates to the process for producing a barrier film, which involves the step of

introducing said auxiliary reductive gas together with said feedstock gas and said nitrogen-containing gas into said vacuum atmosphere.

[0013] The present invention relates to the process for producing a barrier film, which involves the step of introducing said feedstock gas and said nitrogen-containing gas into said vacuum atmosphere without introducing said auxiliary reductive gas.

[0014] The present invention relates to the process for producing a barrier film, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas, said reductive nitrogen-containing gas is introduced at a flow rate once or more higher than the flow rate of said feedstock gas, and said auxiliary reductive gas is introduced at a flow rate once or more but not more than 10 times higher than the flow rate of said reductive nitrogen-containing gas.

[0015] The present invention relates to the process for producing a barrier film, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas, said reductive nitrogen-containing gas is introduced at a flow rate once or more but not more than 5 times higher than the flow rate of said feedstock gas, and said auxiliary reductive gas is introduced at a flow rate 2 times or more but not more than 10 times higher than the flow rate of said reductive nitrogen-containing gas.

[0016] The present invention relates to the process for producing a barrier film, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas, said auxiliary reductive gas is introduced at a flow rate once or more but not more than 15 times higher than the flow rate of the feedstock gas having said high temperature-melting point metal.

[0017] The present invention relates to the process for producing a barrier film, wherein, in the step of growing the thin film of the nitride of said high temperature-melting point metal, a diluent gas not reacting with said high temperature-melting point metal and a gas having an oxygen atom in its chemical structure are introduced so that the pressure of said vacuum atmosphere is regulated to 1 Pa or more but not more than 100 Pa.

[0018] The present invention relates to a process for producing a barrier film which comprising the steps of forming a barrier film made of a thin nitride film of a high temperature-melting point metal on a substrate, wherein the surface of said substrate is exposed to a plasma of hydrogen and a plasma of at least one gas selected from among argon, nitrogen and helium gases and then the thin film of the nitride of said high temperature-melting point metal is formed on the surface of the substrate.

[0019] The invention relates to a barrier film having a thin film of the nitride of a high temperature-melting point metal, wherein said thin nitride film has a content

of said high temperature-melting point metal exceeding the stoichiometric composition ratio thereof.

[0020] The invention relates to a barrier film having a thin nitride film a high temperature-melting point metal formed on a substrate and aiming at preventing the diffusion of metals in a thin interconnecting film formed on said thin nitride film, wherein said thin nitride film is free from silicon.

BRIEF DESCRIPTION OF THE DRAWINGS

[0021]

Figs. 1(a) to (e) show the first half of a flow chart illustrating the process of the present invention.

Figs. 2(f) to (h) show the second half of a flow chart illustrating the process of the present invention.

Figs. 3(i) to (j) show the latter half of a flow chart illustrating the process of the present invention.

Fig. 4 shows an example of a CVD apparatus by which the process of the present invention can be embodied.

Fig. 5 is a graph which shows relationships between the resistivity of a thin tungsten nitride film formed by the process of the present invention and the flow rates of the feedstock gas, the reductive nitrogen-containing gas and the auxiliary reductive gas.

Fig. 6 is a graph which shows a change in the elemental content of a thin tungsten nitride film formed by the process of the present invention depending on the SiH_4/WF_6 flow rate.

Fig. 7 is a graph which shows the composition of tungsten nitride formed by the process of the present invention in the depth direction.

Fig. 8 is a graph which shows the composition of tungsten nitride of the conventional art in the depth direction.

[0022] In these drawings, each numerical symbol has the following meaning:

20	: substrate
24,33	: thin nitride films (barrier films)
25	: thin metal film
27,35	: interconnecting films.

DETAILED DESCRIPTION OF THE INVENTION

[0023] In the present invention which has the above-described constitution, a feedstock gas having a high temperature-melting point metal atom and a reductive nitrogen-containing gas are introduced into a vacuum atmosphere and the feedstock gas is thus reduced by the reductive nitrogen-containing gas to deposit the nitride of the high temperature-melting point metal, wherein a nitrogen-free auxiliary reductive gas is also introduced into the vacuum atmosphere so as to allow

the deposition of the high temperature-melting point metal too.

[0024] Further, a diluent gas and a gas having an oxygen atom in its structure (oxygen gas, etc.) are simultaneously introduced. As a result, it becomes possible to form a nitride film having improved barrier performance and further lowered resistivity, compared with a case where no oxygen atom-containing gas is introduced.

[0025] When the nitride of the high temperature-melting point metal is deposited at a low temperature, the high temperature-melting point metal in the thin nitride film becomes deficient. However, the high temperature-melting point metal atoms deposited by the auxiliary reductive gas compensate for the deficiency. Thus, the obtained thin nitride film has a composition wherein the content of the high temperature-melting point metal exceeds the stoichiometric level thereof.

[0026] The amount of the deposited metal may be smaller than the amount of the deposited nitride. However, the auxiliary reductive gas should be introduced in a somewhat larger amount than the deposition level, since the auxiliary reductive gas is inferior in the reactivity to the reductive nitrogen-containing gas.

[0027] When the auxiliary reductive gas is introduced at an excessively high rate, on the other hand, the content of the high temperature-melting point metal becomes unnecessarily large and, as a result, the thin film exhibits properties close to the properties of the high temperature-melting point metal rather than those of the nitride. When an auxiliary reductive gas containing Si is employed, there arises a problem that the content of Si in the thin nitride film is increased. Therefore, the reductive nitrogen-containing gas, the auxiliary reductive gas, the diluent gas and the gas having an oxygen atom in its structure should be introduced each at an adequate rate.

[0028] For example, ammonia gas (i.e., the reductive nitrogen-containing gas) is introduced at rates 1.0, 2.6 and 5.0 times higher than that of tungsten hexafluoride (i.e., the feedstock gas) and the ratio of silane gas (i.e., the auxiliary reductive gas) to the ammonia gas is varied.

[0029] Fig. 5 shows the results wherein the abscissa indicates the introduction rate of the silane gas expressed by referring the introduction rate of the ammonia gas as to 1 while the ordinate indicates the resistivity of the thin tungsten nitride film thus formed.

[0030] As this graph shows, it is preferable that the reductive nitrogen-containing gas is introduced at a rate amount once or more larger than the introduction rate of the feedstock gas, while the auxiliary reductive gas is introduced at a rate 2 times or more but not more than 10 times larger than the introduction rate of the reductive nitrogen-containing gas.

[0031] The practically available resistivity of a barrier film ranges from about 200 to 300 $\mu\Omega \cdot \text{cm}$. Thus, it can be understood from this graph that the reductive

nitrogen-containing gas is introduced preferably at a flow rate 1 time or more but not more than 5 times higher than the flow rate of the feedstock gas and the auxiliary reductive gas is introduced preferably at a flow rate 2 times or more but not more than 5 times higher than the flow rate of the reductive nitrogen-containing gas.

[0032] Fig. 6 shows the elemental content of each high temperature-melting point metal nitride at various ratios of the flow rate of the feedstock gas to the flow rate of the auxiliary reductive gas in the step of forming high temperature-melting point metal nitrides. In this case, WF_6 is used as the feedstock gas and SiH_4 is used as the auxiliary reductive gas. Oxygen gas is introduced at 1.5 sccm.

[0033] As this graph shows, the high temperature-melting point metal nitride (WN in this case) contains Si when the ratio SiH_4/WF_6 attains 15 or more. This graph also shows that W_xN thin films are formed within the scope of the ratio SiH_4/WF_6 of 1 to 15 time.

[0034] Fig. 7 shows the results of Auger spectrochemical analysis of a thin tungsten nitride film formed at a film forming temperature of 400°C wherein the sputtering time given in the abscissa indicates the depth from the surface. As this graph shows, much tungsten is contained therein (about 4.0 tungsten atoms per nitrogen atom), thereby indicating the effect of the introduction of the auxiliary reductive gas.

[0035] Since the barrier film is oxidized when it is taken out into the atmosphere, oxygen is observed on the film surface. Although silicon is seemingly contained in the film, the content thereof is less than the detection limit, i.e., a measurement error.

[0036] Thus, the resistivity can be lowered by using the high temperature-melting point metal in the thin nitride film at a level exceeding the stoichiometric composition ratio thereof while sustaining the barrier performance.

[0037] In the CMP process, a thin nitride film should be adhered closely to the surface of a substrate. When the surface of the substrate is cleaned with a plasma of at least one gas selected from among argon, nitrogen and helium gases, the plasma and the plasma of hydrogen gas, and a plasma of mixed nitrogen, the thin nitride film formed on this surface would not peel off even under a load of 1 kg/cm^2 , thereby achieving an adhesiveness usable in the CMP process.

[0038] When the thin nitride film contains silicon, the silicon reacts with the high temperature-melting point metal (tungsten, etc.) at a high temperature to form a silicon compound such as tungsten silicide to thereby increase the resistivity. Since the thin nitride film of the present invention is free from silicon, no silicon compound is formed and thus the resistivity can be stabilized at a low level.

[0039] For comparison, Fig. 8 shows the results of Auger spectrochemical analysis of a thin tungsten nitride film formed by the CVD method of the conven-

tional art at a film forming temperature of 400°C. This thin film contains about 1.7 tungsten atoms per nitrogen atom, i.e., showing a smaller tungsten content. It also shows a high resistivity of 1000 $\mu\Omega\text{cm}$ or more.

[0040] In the step of the formation of the thin nitride film of the high temperature-melting point metal, it is preferable to regulate the pressure to 1 Pa or more but not more than 10000 Pa, still preferably 1 Pa or more but not more than 100 Pa.

DESCRIPTION OF THE EMBODIMENTS OF THE INVENTION

[0041] Now, embodiments of the present invention will be described by reference to the attached drawings.

[0042] Figs. 1(a) to (d) show a flow chart which shows an example of the embodiment of the present invention. In Fig. 1(a), the numerical symbol 20 shows a substrate to be treated. This substrate 20 has a semiconductor substrate 21 made of single silicon crystal on the surface of which a primary coat film 22 and an insulation film 23 made of silicon oxide are formed. The primary film 22 and the insulation film 23 are provided with a hole 31. The surface of semiconductor substrate 21 is exposed at a bottom 32 of the hole 31.

[0043] A barrier film is to be formed on the surface of this substrate 20.

[0044] In Fig. 4, the numerical symbol 50 shows a CVD apparatus whereby the present invention can be embodied. This CVD apparatus 50 has a vacuum chamber 51 which is connected to a transport room (not shown). The vacuum chamber 51 is provided with a substrate holder 53 in the bottom side and an electrode 55 in the top side.

[0045] When a barrier film is to be formed on the substrate 20 by using this CVD apparatus 50, the substrate 20 is first brought into the transport room and the transport room and the vacuum chamber 51 are evacuated. Next, a gate valve 52 between the vacuum chamber 51 and the transport room is opened and then the substrate 20 is brought into the CVD apparatus 50.

[0046] The substrate holder 53 is provided with a substrate elevator unit 54 with which the substrate brought into the vacuum chamber 51 is placed on the substrate holder 53. In Fig. 4, the substrate 20 is thus placed on the substrate holder 53.

[0047] Subsequently, the heater in the substrate holder 53 is switched on and thus the substrate 20 is heated to a temperature of not lower than 300°C but not higher than 400°C.

[0048] The vacuum chamber 51 is provided with a gas inlet system 57 through which argon gas and ammonia gas are introduced into the vacuum chamber 51 each at a prescribed flow rate. By applying a high-frequency voltage between the substrate holder 53 and the electrode 55, ionized nitrogen and hydrogen are liberated from the ammonia gas. In this step, the ionized argon gas serving as a diluent gas is mixed with the ion-

ized nitrogen and hydrogen to give a plasma.

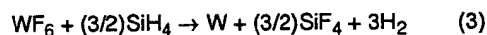
[0049] The insulation film 23 on the surface of the substrate 20 is faced closely to the electrode 55. As the plasma is formed, the surface of the insulation film 23 and the surface of the semiconductor substrate 21 at the bottom of the hole 31 are exposed to the plasma and the organic deposits are decomposed (i.e., cleaning).

[0050] The cleaning is performed at an ammonia gas flow rate of 75 sccm, an argon gas flow rate of 240 sccm, under a pressure of 40 Pa and at a high-frequency electrical power of 100 W. After cleaning for about 50 seconds, the application of the high-frequency voltage is stopped to thereby extinguish the plasma. Although argon (Ar) gas is used in this case, it can be substituted by nitrogen (N_2) gas, helium (He) gas or a mixture thereof.

[0051] Next, the flow rates of the ammonia gas and the argon gas are changed and, at the same time, tungsten hexafluoride (WF_6) gas, silane gas and oxygen gas are introduced into the vacuum chamber 51 through the gas inlet system 57 together with the ammonia gas and the argon gas.

[0052] Since the ammonia gas is superior in the reactivity to the silane gas, the tungsten hexafluoride serves as the feedstock gas and the ammonia gas serves as the reductive nitrogen-containing gas. Thus, the reaction of reducing the feedstock gas proceeds. Since the ammonia gas contains nitrogen, tungsten nitride is deposited on the surface of the insulation film 23 and on the surface of the semiconductor substrate 21 in the hole 31 due to the reduction reaction represented by the above formula (1).

[0053] Although the silane gas introduced into the vacuum chamber 51 also has reductive properties, it is inferior in reactivity to the ammonia gas and, therefore, serves as the auxiliary reductive gas. Since the silane gas has no nitrogen atom, the feedstock gas is reduced thereby in accordance with the following formula (3) and thus metal tungsten is deposited.



[0054] When metal tungsten is deposited, it is incorporated into the thin tungsten nitride film under growing. That is to say, metal tungsten is supplied during the growth of the thin tungsten nitride film. When the thin tungsten nitride film grows at a low temperature, the deficiency of tungsten is thus compensated for and a barrier film (a thin tungsten nitride film) having a composition close to the stoichiometric composition or one having a high temperature-melting point metal content exceeding the stoichiometric level can be formed.

[0055] The thin tungsten nitride film may grow under, for example, the following conditions; substrate temperature: 380°C; feedstock gas flow rate: 5 sccm; reductive nitrogen-containing gas flow rate: 13 sccm; auxiliary reductive gas flow rate: 39 sccm; argon gas

flow rate: 240 sccm; oxygen gas introduction rate: 1.5 sccm; pressure: 40 Pa.

[0056] When the feedstock gas is subjected to the reduction reaction by the reductive nitrogen-containing gas and the auxiliary reductive gas for a prescribed period of time, a thin tungsten nitride film is formed on the surface of insulation film 23 and on the surface of the semiconductor substrate 21, as shown in Fig. 1(b)24.

[0057] Next, the supply of the reductive nitrogen-containing gas is ceased and the flow rate of the auxiliary reductive gas is increased. Thus, the feedstock gas is reduced by the auxiliary reductive gas and metal tungsten is deposited. In Fig. 1 (c), the numerical symbol 25 shows a thin metal tungsten film growing on the surface of the thin nitride film 24.

[0058] The thin metal tungsten film may be formed under, for example, the following conditions; substrate temperature: 380°C; feedstock gas flow rate: 20 sccm; auxiliary reductive gas flow rate: 5 sccm; diluent (argon) gas introduction rate: 240 sccm; pressure: 40 Pa.

[0059] Although the thin nitride film 24 has a favorable barrier performance against copper, it has a higher resistivity than that of the high temperature-melting point metal. On the other hand, the film of the high temperature-melting point metal (for example, thin metal tungsten film, etc.) has a poor barrier performance against copper but has a much lower resistivity than that of the thin nitride film 24.

[0060] When a thin nitride film 24 is formed as a barrier film and a thin high temperature-melting point metal film is laminated thereon as in the above-described case, the resistivity can be lowered while sustaining the favorable barrier performance against copper.

[0061] On the contrary, a thin nitride film may be formed on a high temperature-melting point metal film or the thin nitride film may be formed as a single layer.

[0062] After allowing the thin tungsten film 25 to grow for 20 to 30 seconds under the above-described conditions, the substrate 20 is taken out from the CVD apparatus 50. Then a thin copper film is grown on the surface of the thin high temperature-melting point metal film 25 by the metal plating method, the sputtering method, etc. In Fig. 1(d), the numerical symbol 26 shows this thin copper film 26.

[0063] After the formation of the thin copper film 26, the surface is abraded by the CMP method to thereby eliminate the thin copper film 26, the thin nitride film 24 and the thin metal film 25 on the insulation film 23. Thus, an interconnecting film 27 made of the thin copper film 26 is formed in the hole 31. The thin nitride film 24 is located between the interconnecting film 27 and the semiconductor substrate 21 and between the interconnecting film 27 and the insulation film 23 so as to prevent copper from diffusion.

[0064] Next, primary films 41 and 43 and insulation films 42 and 44 are laminated alternately and the sur-

face of the interconnecting film 27 is windowed to form a hole or a groove, as shown in Fig. 2(g). In Fig. 2(g), the numerical symbol 38 shows the groove or hole. The interconnecting film 27 is exposed on the bottom of this groove or hole 38.

[0065] Then the substrate 20 is brought into the CVD apparatus 50 and a thin nitride film (a thin tungsten nitride film) is formed under the same conditions as in forming the thin nitride film shown in Fig. 1(b).

[0066] The numerical symbol 33 in Fig. 2(h) shows the thus formed thin nitride film which covers the inner face of the groove or hole 38 and the surface of the insulation film 44 and the surface of the interconnecting film 27.

[0067] Subsequently, the thin copper film 34 is grown by the metal plating method or the sputtering method, as shown in Fig. 3(i). Thus, the groove or hole 38 is filled with the thin copper film 34.

[0068] Finally, the surface is abraded by the CMP method. Thus, an interconnecting film 35 is formed by the thin copper film 34 filled into the groove or hole 38, as shown in Fig. 3(j).

[0069] Since the barrier film 33 is located between the interconnecting film 35 and the insulation films 42 and 44, the diffusion of copper into the insulation films 42 and 44 can be prevented.

[0070] In the above-described case, a thin tungsten nitride film is formed by using tungsten as the high temperature-melting point metal, ammonia gas as the reductive nitrogen-containing gas and silane gas as the auxiliary reductive gas. Although tungsten hexafluoride is used as the feedstock gas in the above case, it is also possible to use $W(CO)_6$ gas therefor.

[0071] Moreover, the present invention involves in its scope cases where barrier films are produced by using high temperature-melting point metals other than tungsten and forming thin films of the nitrides thereof. When titanium (Ti) is used as the high temperature-melting point metal, titanium halide gases (TiF_4 , $TiCl_4$, etc.) are usable as the feedstock gas. When tantalum (Ta) is used as the high temperature-melting point metal, tantalum halide gases ($TaCl_5$, etc.) are usable as the feedstock gas. It is also possible to use halides of Mo or Nb as the feedstock gas.

[0072] As the reductive nitrogen-containing gas containing a nitrogen atom, use can be made of N_2H_4 gas, NF_3 gas, N_2O gas, etc. in addition to NH_3 gas.

[0073] As the nitrogen-free auxiliary reductive gas, use can be made of H_2 gas, Si_2H_6 gas, PH_3 gas, B_2H_6 gas etc. in addition to SiH_4 gas.

[0074] As the diluent gas, it is also possible to use argon gas, nitrogen gas, helium gas or a mixture of these gases.

[0075] The cleaning conditions as described above are given merely by way of example. Therefore, the cleaning may be performed under different conditions. For example, similar effects can be achieved by carrying out the cleaning for 60 seconds at an argon gas flow

rate of 100 sccm, under a pressure of 1.0 Pa at a high-frequency electric power of 150W.

[0076] After completing the cleaning under such conditions, additional cleaning may be performed by adding ammonia gas to the argon gas.

[0077] According to the present invention, a barrier film (a thin film of the nitride of a high temperature-melting point metal) having a low resistance can be formed by the CVD method at a temperature of 500°C or lower, in particular, from 350 to 450°C. Therefore, interlayer insulation films are not damaged thereby.

[0078] Furthermore, the thin nitride film is formed by the heat CVD method, which makes it possible to achieve good step coverage.

Claims

1. A process for producing a barrier film which comprises the steps of:

providing a substrate in a vacuum atmosphere; introducing a feedstock gas having a high temperature-melting point metal in its structure and a reductive nitrogen-containing gas comprising a nitrogen atom into said vacuum atmosphere; and

forming a thin film of the nitride of said high temperature-melting point metal on said substrate;

wherein a nitrogen-free auxiliary reductive gas is introduced into said vacuum atmosphere.

2. The process for producing a barrier film according to claim 1, which comprises a step of introducing said auxiliary reductive gas together with said feedstock gas and said reductive nitrogen-containing gas into said vacuum atmosphere.

3. The process for producing a barrier film according to claim 2, which comprises a step of introducing said feedstock gas and said reductive nitrogen-containing gas into said vacuum atmosphere without introducing said auxiliary reductive gas.

4. The process for producing a barrier film according to claim 2, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas;

said reductive nitrogen-containing gas is introduced at a flow rate once or more higher than the flow rate of said feedstock gas, and said auxiliary reductive gas is introduced at a flow rate once or more but not more than 10 times higher than the flow rate of said reductive nitrogen-containing gas.

5. The process for producing a barrier film according

to claim 1, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas;

said reductive nitrogen-containing gas is introduced at a flow rate once or more but not more than 5 times higher than the flow rate of said feedstock gas, and said auxiliary reductive gas is introduced at a flow rate 2 times or more but not more than 10 times higher than the flow rate of said reductive nitrogen-containing gas.

6. The process for producing a barrier film according to claim 2, wherein, in the step of introducing said auxiliary reductive gas together with said reductive nitrogen-containing gas and said feedstock gas;

said auxiliary reductive gas is introduced at a flow rate once or more but not more than 15 times higher than the flow rate of the feedstock gas having said high temperature-melting point metal.

7. The process for producing a barrier film according to claim 1, wherein, in the step of growing the thin film of the nitride of said high temperature-melting point metal;

a diluent gas not reacting with said high temperature-melting point metal and a gas having an oxygen atom in its chemical structure are introduced so that the pressure of said vacuum atmosphere is regulated to 1 Pa or more but not more than 100 Pa.

8. A process for producing a barrier film for forming a barrier film made of a thin film of the nitride of a high temperature-melting point metal on a substrate, wherein;

the surface of said substrate is exposed to a plasma of hydrogen gas and a plasma containing at least one gas selected from among argon, nitrogen and helium gases, and then the thin film of the nitride of said high temperature-melting point metal is formed on the surface of the substrate.

9. A barrier film comprising a thin nitride film of a high temperature-melting point metal, wherein;

said thin nitride film has a content of said high temperature-melting point metal exceeding the stoichiometric composition ratio thereof.

10. A barrier film comprising a thin nitride film of a high temperature-melting point metal formed on a substrate and aiming at preventing the diffusion of met-

als in an interconnecting thin film formed on said thin nitride film, wherein;

said thin nitride film is free from silicon.

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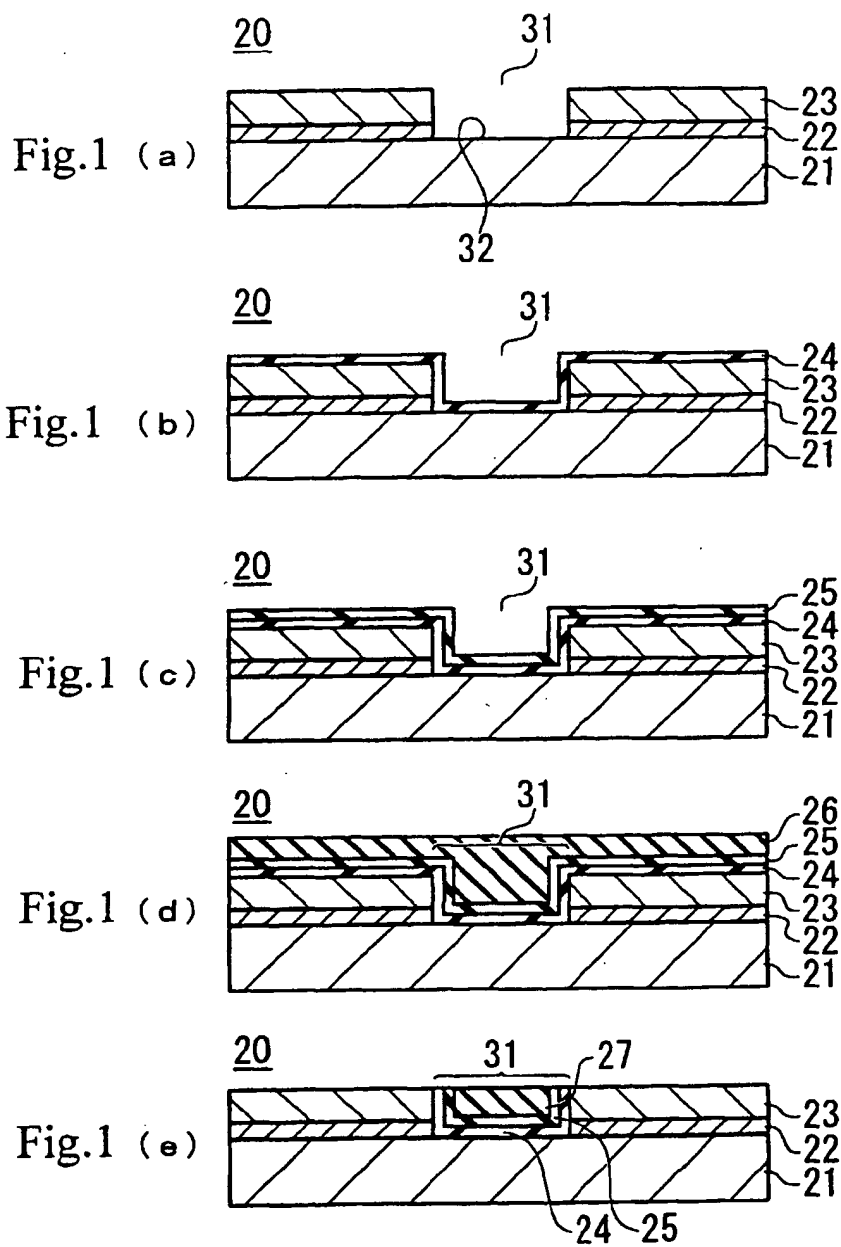
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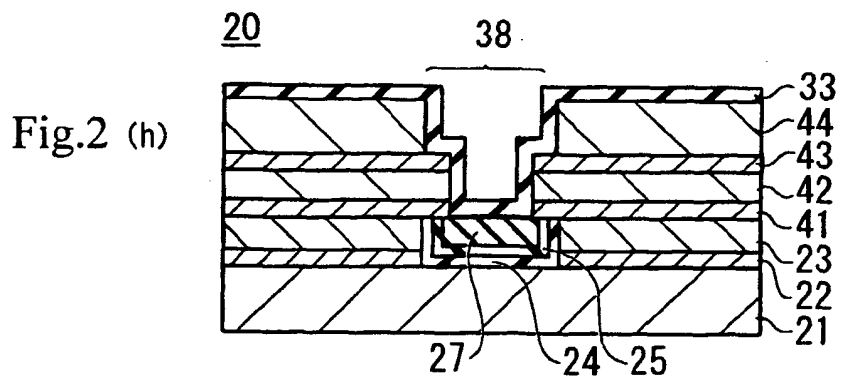
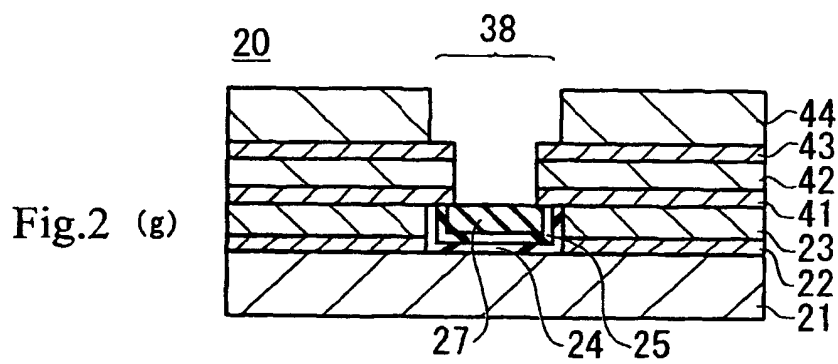
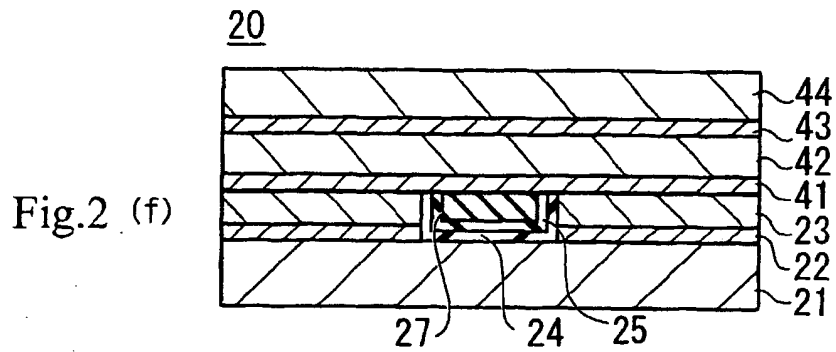
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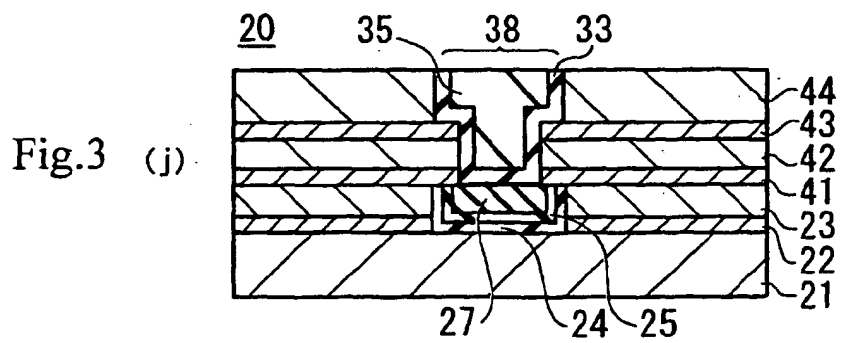
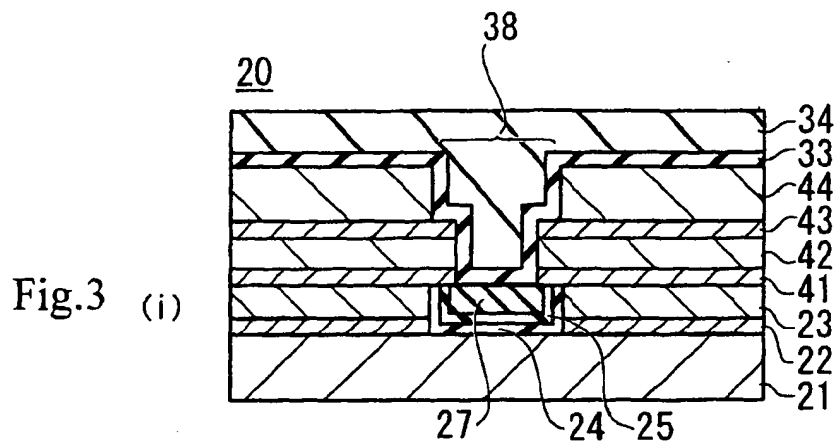


Fig.4

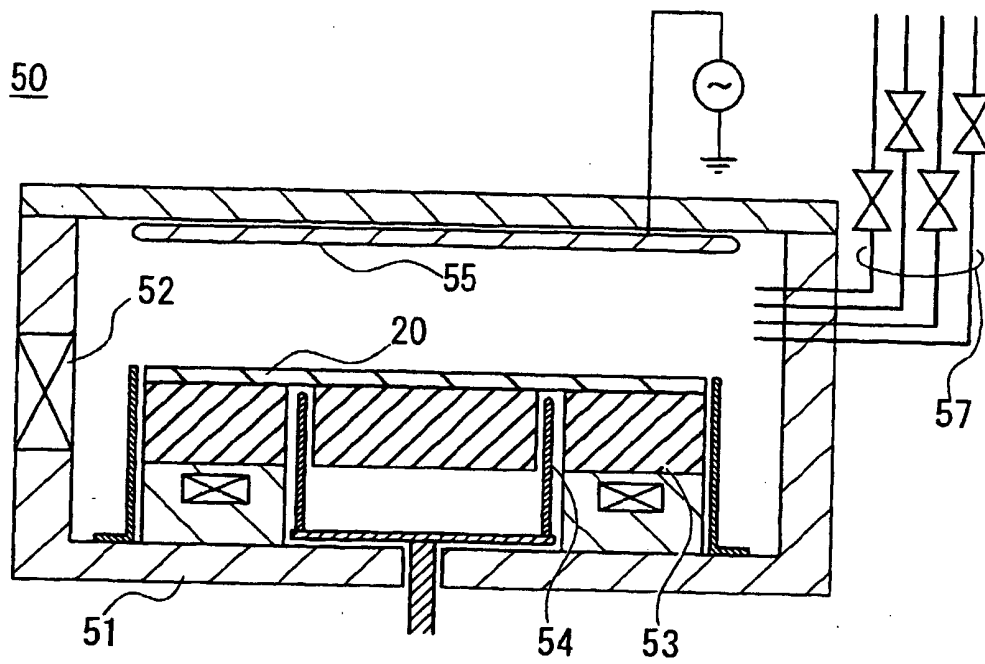


Fig.5

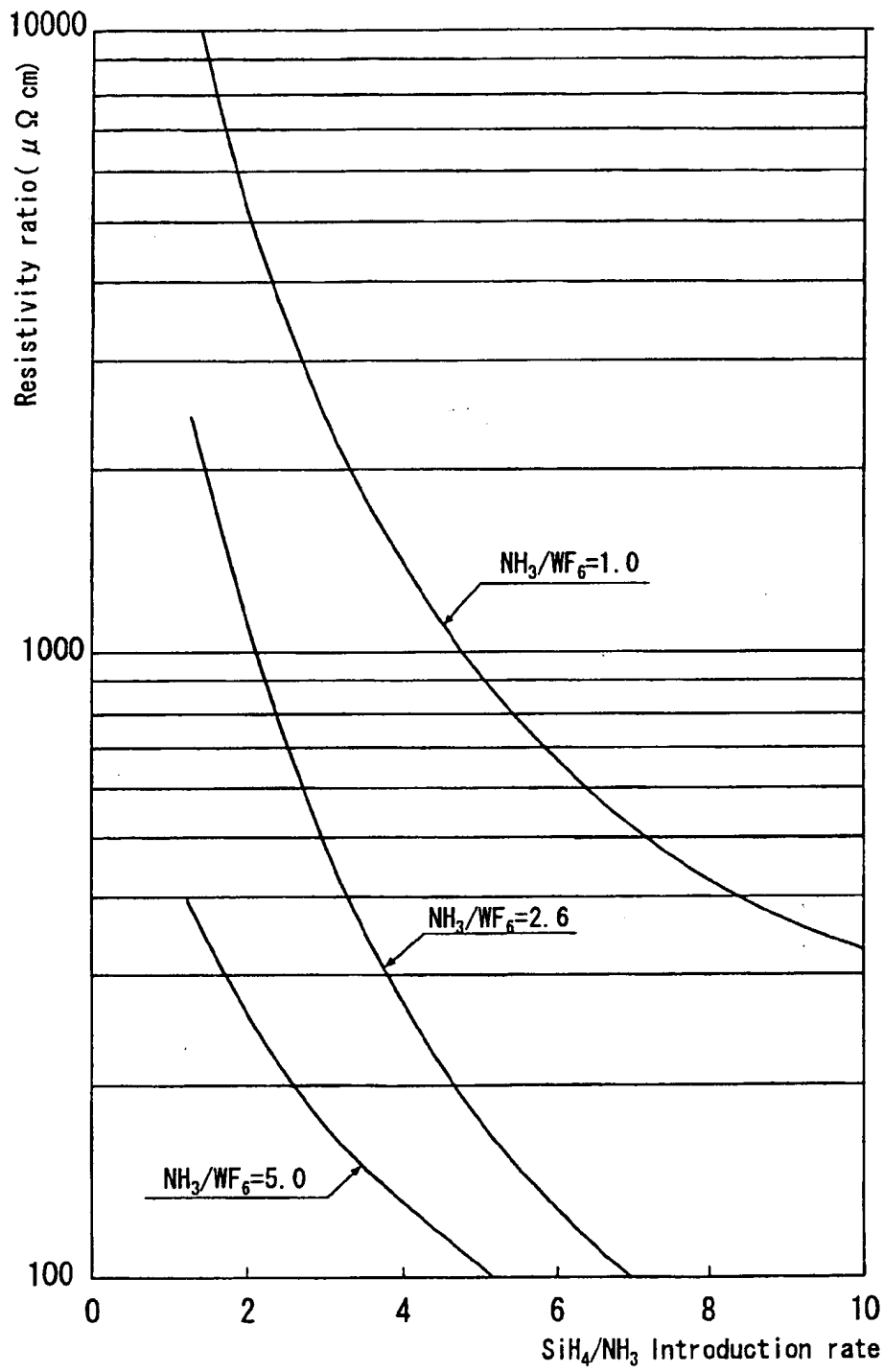


Fig.6

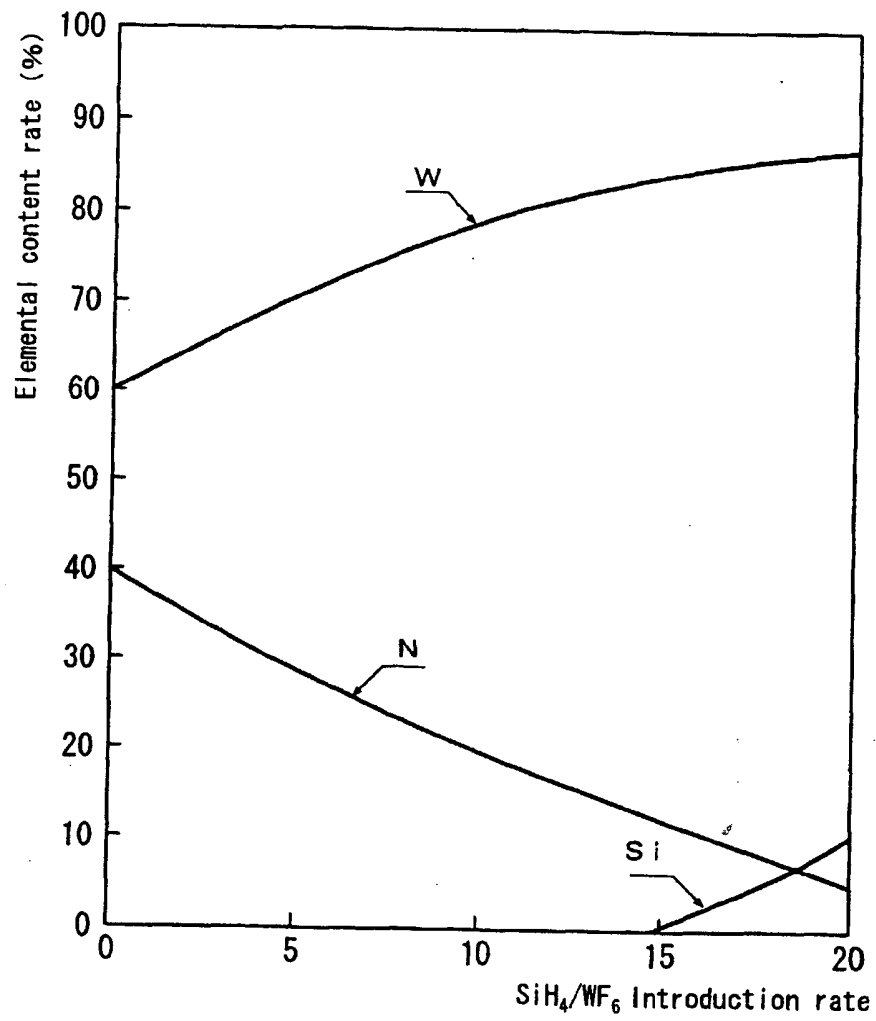


Fig.7

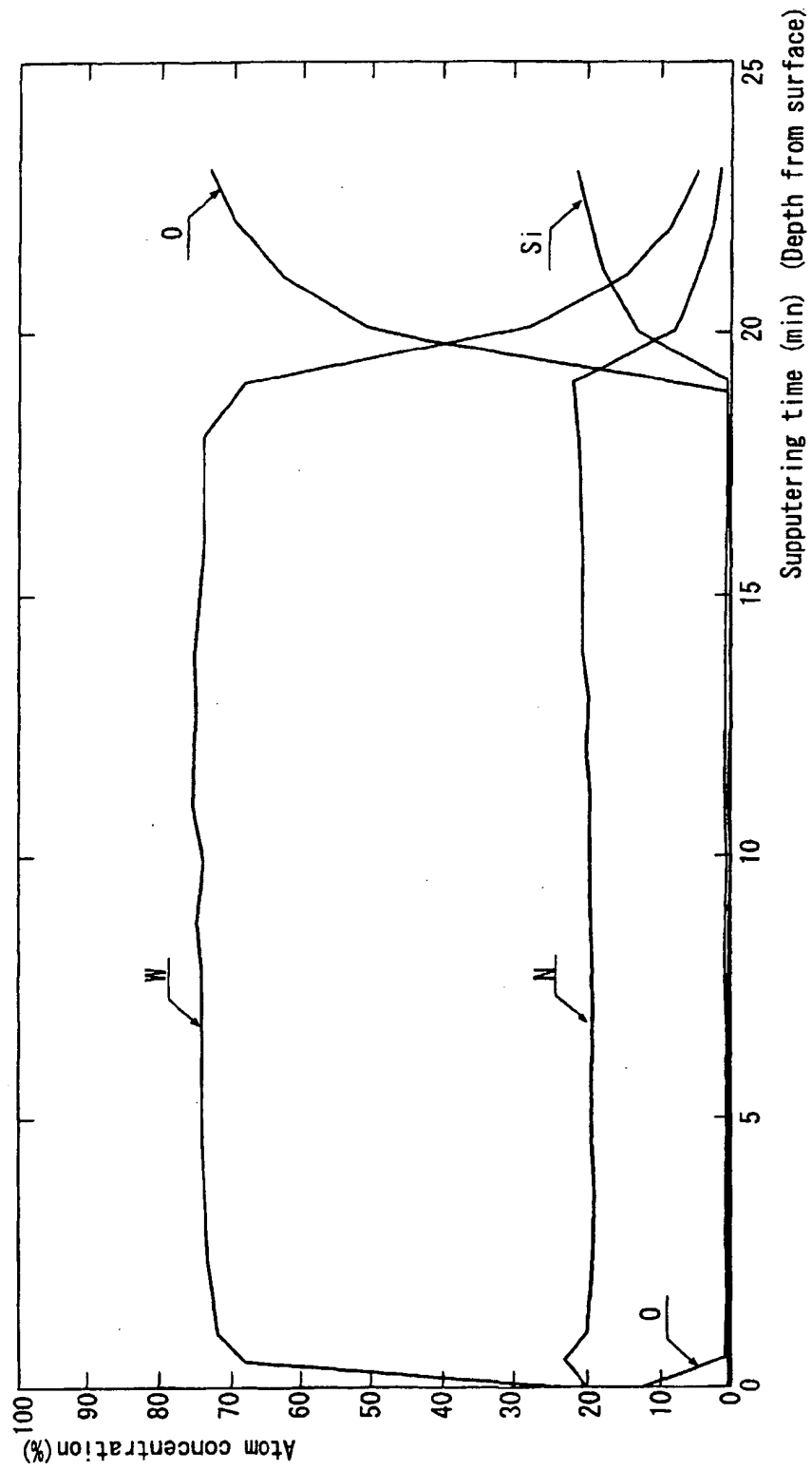
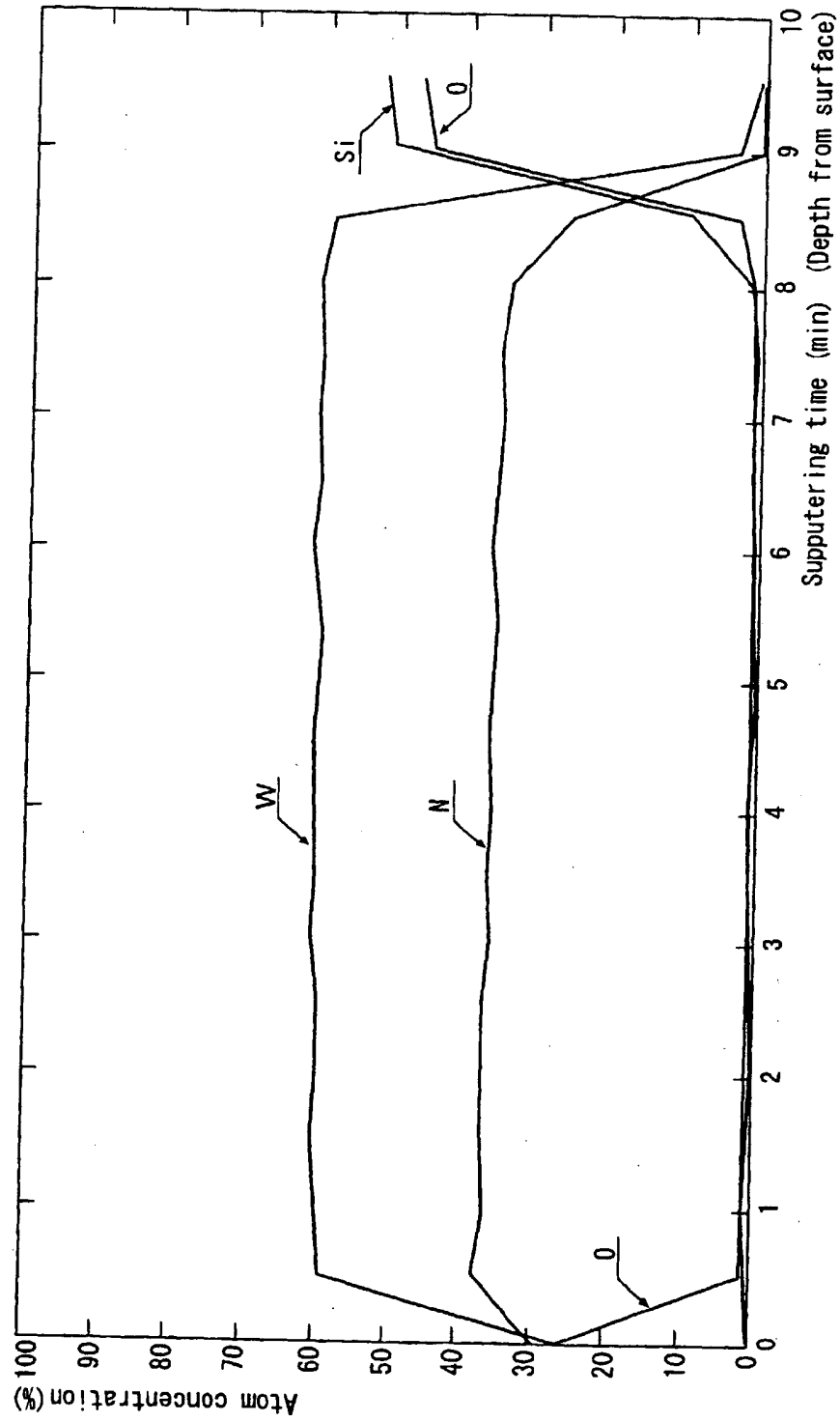


Fig.8 Prior Art





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EUROPEAN SEARCH REPORT

Application Number
EP 00 10 3073

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	PATENT ABSTRACTS OF JAPAN vol. 1995, no. 01, 28 February 1995 (1995-02-28) & JP 06 283532 A (KAWASAKI STEEL CORP), 7 October 1994 (1994-10-07)	1,2,6,10	C23C16/34 C23C16/30 H01L21/285
Y	* abstract *	3	
X	WO 98 33950 A (TOKYO ELECTRON LTD ;TOKYO ELECTRON ARIZONA INC (US)) 6 August 1998 (1998-08-06) * page 12, line 6 - page 13, line 15 *	1,4,5	
X	WO 96 17104 A (MICRON TECHNOLOGY INC) 6 June 1996 (1996-06-06)	1,6	
Y	* page 4, line 13 - page 5, line 4 *	7	
X	EP 0 536 664 A (SUMITOMO METAL IND) 14 April 1993 (1993-04-14) * page 4, line 41 - line 49 *	9,10	
X	EP 0 840 363 A (TEXAS INSTRUMENTS INC) 6 May 1998 (1998-05-06) * figure 1 *	8	TECHNICAL FIELDS SEARCHED (Int.Cl.7) C23C H01L
Y	LEE J ET AL: "MOCVD OF TIN AND/OR TI FROM NEW PRECURSORS" THIN SOLID FILMS,CH,ELSEVIER-SEQUOIA S.A. LAUSANNE, vol. 320, no. 1, 4 May 1998 (1998-05-04), pages 15-19, XP000668459 ISSN: 0040-6090 * table 1 *	3	
Y	DE 35 22 427 A (FISCHER HELMUT DIPL ING;MUELLER JOERG PROF DR ING) 20 February 1986 (1986-02-20) * page 2, line 5 - line 13 * * page 11, line 4 - line 18 *	7	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 16 May 2000	Examiner Ekhult, H
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

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**LACK OF UNITY OF INVENTION
SHEET B**

Application Number
EP 00 10 3073

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

1. Claims: 1-7,9,10

Process for deposition of a nitride film.

2. Claim : 8

Plasma pretreatment process.



European Patent
Office

Application Number

EP 00 10 3073

CLAIMS INCURRING FEES

The present European patent application comprised at the time of filing more than ten claims.

- ☐ Only part of the claims have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims and for those claims for which claims fees have been paid, namely claim(s):
- ☐ No claims fees have been paid within the prescribed time limit. The present European search report has been drawn up for the first ten claims.

LACK OF UNITY OF INVENTION

The Search Division considers that the present European patent application does not comply with the requirements of unity of invention and relates to several inventions or groups of inventions, namely:

see sheet B

- ☐ All further search fees have been paid within the fixed time limit. The present European search report has been drawn up for all claims.
- ☒ As all searchable claims could be searched without effort justifying an additional fee, the Search Division did not invite payment of any additional fee.
- ☐ Only part of the further search fees have been paid within the fixed time limit. The present European search report has been drawn up for those parts of the European patent application which relate to the inventions in respect of which search fees have been paid, namely claims:
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**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 10 3073

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16-05-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
JP 06283532 A	07-10-1994	NONE	
WO 9833950 A	06-08-1998	US 5989652 A AU 5917698 A DE 19882045 T GB 2337529 A	23-11-1999 25-08-1998 23-12-1999 24-11-1999
WO 9617104 A	06-06-1996	AU 4290396 A EP 0742847 A EP 0982772 A JP 9509288 T US 5691235 A	19-06-1996 20-11-1996 01-03-2000 16-09-1997 25-11-1997
EP 0536664 A	14-04-1993	DE 69216747 D DE 69216747 T JP 5211134 A US 5508066 A	27-02-1997 31-07-1997 20-08-1993 16-04-1996
EP 0840363 A	06-05-1998	JP 10144627 A	29-05-1998
DE 3522427 A	20-02-1986	NONE	

EPO FORM P4459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

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